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L3: Entry 30 of 30

File: USPT

Dec 22, 1998

DOCUMENT-IDENTIFIER: US 5852035 A

TITLE: Method for inhibiting neoplastic cells and related conditions by exposure to substituted N- arylmethyl and heterocyclmethyl-1H-pyrazolo (3,4-B) quinolin-4-amines

Application Filing Date (1):19971212INVENTOR (2):Piazza; Gary A.Brief Summary Text (4):

Each year in the United States alone, untold numbers of people develop precancerous lesions, which is a form of neoplasia, as discussed below. Such lesions exhibit a strong tendency to develop into malignant tumors, or cancer. Such lesions include lesions of the breast (that can develop into breast cancer), lesions of the skin (that can develop into malignant melanoma or basal cell carcinoma), colonic adenomatous polyps (that can develop into colon cancer), and other such neoplasms. Compounds that prevent or induce the remission of existing precancerous or cancerous lesions or carcinomas would greatly reduce illness and death from cancer.

Brief Summary Text (8):

In view of these grim statistics, efforts in recent years have concentrated on colon cancer prevention. Colon cancer usually arises from pre-existing benign neoplastic growths known as polyps. Prevention efforts have emphasized the identification and removal of colonic polyps. Polyps are identified by x-ray and/or colonoscopy, and usually removed by devices associated with the colonoscope. The increased use of colon x-rays and colonoscopies in recent years has detected clinically significant precancerous polyps in four to six times the number of individuals per year that acquire colon cancer. During the past five years alone, an estimated 3.5 to 5.5 million people in the United States have been diagnosed with adenomatous colonic polyps, and it is estimated that many more people have or are susceptible to developing this condition, but are as yet undiagnosed. In fact, there are estimates that 10-12 percent of people over the age of 40 will form clinically significant adenomatous polyps.

Brief Summary Text (9):

Removal of polyps has been accomplished either with surgery or fiber-optic endoscopic polypectomy--procedures that are uncomfortable, costly (the cost of a single polypectomy ranges between \$1,000 and \$1,500 for endoscopic treatment and more for surgery), and involve a small but significant risk of colon perforation. Overall, about \$2.5 billion is spent annually in the United States in colon cancer treatment and prevention.

Brief Summary Text (12):

In most cases (i.e. the cases of sporadic lesion formation, e.g. so-called common sporadic polyps), lesion removal will be effective to reduce the risk of cancer. In a small percentage of cases (i.e. cases where numerous lesions form, e.g. the so-called polyposis syndromes), removal of all or part of the effected area (e.g. the colon) is indicated. For example, the difference between common sporadic polyps and polyposis syndromes is dramatic. Common sporadic polyp cases are characterized by relatively few polyps which can usually be removed leaving the colon intact. By contrast, polyposis syndrome cases can be characterized by many (e.g. hundreds or more) of polyps--literally covering the colon in some cases--making safe removal of the polyps impossible short of surgical removal of the colon.

Brief Summary Text (18):

In recent years, several non-steroidal anti-inflammatory drugs ("NSAIDs"), originally developed to treat arthritis, have shown effectiveness in inhibiting and eliminating colonic polyps. Polyps virtually disappear when the patients take the drug, particularly when the NSAID sulindac is administered. However, the prophylactic use of currently available NSAIDs, even in polyposis syndrome patients, is marked by severe side reactions that include gastrointestinal irritations and ulcerations. Once NSAID treatment is terminated due to such complications, the polyps return, particularly in polyposis syndrome patients.

Brief Summary Text (52):

As used herein, the term "precancerous lesion" includes syndromes represented by abnormal neoplastic, including dysplastic, changes of tissue. Examples include adenomatous growths in colonic, breast or lung tissues, or conditions such as dysplastic nevus syndrome, a precursor to malignant melanoma of the skin. Examples also include, in addition to dysplastic nevus syndromes, polyposis syndromes, colonic polyps, precancerous lesions of the cervix (i.e., cervical dysplasia), prostatic dysplasia, bronchial dysplasia, breast, bladder and/or skin and related conditions (e.g., actinic keratosis), whether the lesions are clinically identifiable or not.

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DATE-ISSUED: December 22, 1998

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Pamukcu; Rifat	Spring House	PA		
<u>Piazza</u> ; Gary A.	Doylestown	PA		

ASSIGNEE-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY	TYPE CODE
Cell Pathways, Inc.	Horsham	PA			02

APPL-NO: 08/ 989357 [PALM]

DATE FILED: December 12, 1997

INT-CL: [06] A61 K 31/44, A61 K 31/535

US-CL-ISSUED: 514/293; 514/236.5

US-CL-CURRENT: 514/293; 514/236.5

FIELD-OF-SEARCH: 514/293, 514/236.5

PRIOR-ART-DISCLOSED:

U.S. PATENT DOCUMENTS

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<input type="checkbox"/>	<u>3161654</u>	December 1964	Shen	260/319
<input type="checkbox"/>	<u>3322755</u>	May 1967	Roch et al.	260/246
<input type="checkbox"/>	<u>3517005</u>	June 1970	Cronin et al.	260/256.4
<input type="checkbox"/>	<u>3594480</u>	July 1971	Cronin et al.	424/250
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<input type="checkbox"/>	<u>3654349</u>	April 1972	Shen et al.	260/615M
<input type="checkbox"/>	<u>3780040</u>	December 1973	Schnettler et al.	260/256.5
<input type="checkbox"/>	<u>3812127</u>	May 1974	Cronin et al.	260/268BQ

<input type="checkbox"/>	<u>3819631</u>	June 1974	Broughton et al.	260/256.4F
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<input type="checkbox"/>	<u>4039544</u>	August 1977	Broughton et al.	260/256.4F
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<input type="checkbox"/>	<u>5614627</u>	March 1997	Takase et al.	544/293

FOREIGN PATENT DOCUMENTS

FOREIGN-PAT-NO	PUBN-DATE	COUNTRY	US-CL
0 347146 A2	December 1989	EP	
0 349239 A2	January 1990	EP	
0 351058	January 1990	EP	
0 352960 A2	January 1990	EP	
0 395328 A2	October 1990	EP	
0 428268 A2	May 1991	EP	
0 463756 A1	January 1992	EP	
0 485158 A2	May 1992	EP	
0 485173 A2	May 1992	EP	
0 485172 A2	May 1992	EP	
0 485171 A2	May 1992	EP	
0 485157 A2	May 1992	EP	
0 508586 A1	October 1992	EP	
0 526004 A1	February 1993	EP	
0 607439 A1	July 1994	EP	
3038166	May 1981	DE	
56-53659 A	May 1981	JP	
57-167974 A	October 1982	JP	
807826	January 1959	GB	
2063249	June 1981	GB	
WO 92/03419	March 1992	WO	
WO 93/07149	April 1993	WO	
WO 93/12095	June 1993	WO	
WO 94/05661	March 1994	WO	
WO 97/03985	February 1997	WO	

OTHER PUBLICATIONS

Waddell, W.R. et al., Am. J. Surgery, vol. 157, pp. 175-179 (1989).
 Gonzaga, R.A.F. et al., The Lancet, Mar. 30, 1985, p. 751.
 Waddell, W.R. et al., J. Surg. Oncology, vol. 24, pp. 83-87 (1983).
 Federation Proceedings (1972) of the Federation of American Societies for Experimental Biology abstract Nos. 2044 and 2045.
 Gilman, S.C. et al., Nonsteroidal Anti-inflammatory Drugs in Cancer Therapy, (circa 1985).
 Brogden, R.N. et al., Drugs, vol. 16, pp. 97-114 (1978).
 Hucker, H.B. et al., Drug Metabolism & Disposition, vol. 1, No. 6, pp. 721-736 (1973).
 Shen, T.Y. et al., Chemical and Biological Studies on Indomethacin, Sulindac and Their Analogs, pp. 107-178 (circa 1975).
 Duggan, D.E. et al., Clin. Pharm. & Therapeutics, vol. 21, No. 3, pp. 326-335 (1976).
 Duggan, D.E. et al., J. Pharm. & Exper. Therap., vol. 201, No. 1, pp. 8-13 (1977).
 Glavin, G.B. et al., Toxicology and Applied Pharmacology, vol. 83, pp. 386-389 (1986).
 Moorghen, M. et al., Journal of Pathology, vol. 156, pp. 341-347 (1988).
 Moorghen, M. et al., Acta Histochemica, Suppl.-Band XXIX, S. 195-199 (1990).
 Bjarnason et al., Gastroenterology, vol. 94, No. 4, pp. 1070-1074 (1988).
 Badrieh, Y., et al., Chem. Ber., vol. 125, pp. 667-674 (1992).
 Silvola, J. et al., Effects of nonsteroidal anti-inflammatory drugs on rat gastric mucosal phosphodiesterase activity, Agents and Actions, vol. 12.4, pp. 516-520 (1982).
 Curtis-Prior, P.B. et al., Cyclic Nucleotide Phosphodiesterase Activity of Human Normal and Carcinomatous Lung Tissue, The Lancet, pp. 1225-1225 Dec. 4, 1976.
 Pepin, P. et al., Effects of Sulindac and Oltipraz on the tumorigenicity of 4-(methylnitrosamino) 1-(3-pyridyl)-1-Butanone in A/J mouse lung, Carcinogenesis, vol. 13, No. 3, pp. 341-348 (1992).
 Nicholson, C.D. et al. Differential modulation of tissue function and therapeutic potential of selective inhibitors of cyclic nucleotide phosphodiesterase isoenzymes, Trends Pharmacol. Sci. (TiPS), vol. 12, pp. 19-27 (1991).
 Ahn, H.S. et al., Effects of Selective Inhibitors on Cyclic Nucleotide Phosphodiesterases of Rabbit Aorta, Biochemical Pharmacology, vol. 38, No. 19, pp. 3331-3339 (1989).
 Luginer, C. et al., Selective Inhibition of cyclic Nucleotide Phosphodiesterases of

- Human, Bovine and Rat Aorta, *Biochem. Pharmacology*, vol. 35, No. 10, pp. 1743-1751 (1986).
- Turner, N.C. et al., Relaxation of guinea-pig trachea by cyclic AMP phosphodiesterase inhibitors and their enhancement by sodium nitroprusside, *Br. J. Pharmacol.* vol. III, pp. 1047-1052 (1994).
- Weishaar, R.E. et al., Multiple Molecular Forms of Cyclic Nucleotide Phosphodiesterase in Cardiac and Smooth Muscle and in Platelets, *Biochem. Pharmacology*, vol. 35, No. 5, pp. 787-800 (1986).
- Murray, K.J. et al., Potential Use of Selective Phosphodiesterase Inhibitors in the Treatment of Asthma, *New Drugs for Asthma Therapy*, Birkhauser Verlag Basel, pp. 27-46 (1991).
- Saeki, T. et al., Isolation of Cyclic Nucleotide Phosphodiesterase Isozymes From Pig Aorta, *Biochem. Pharmacology*, vol. 46, No. 5, pp. 833-839 (1993).
- Turner, N.C. et al., Pulmonary effects of type V cyclic GMP specific phosphodiesterase inhibition in anaesthetized guinea-pig, *Br. J. Pharmacol.*, vol. 111, 1198-1204 (1994).
- Ferreira, S.H. et al., The molecular mechanism of action of peripheral morphine analgesia: stimulation of the cGMP system via nitric oxide release, *European Journal of Pharmacology*, 201 pp. 121-122 (1991).
- Hidaka, H. et al., Selective Inhibitors of Three Forms of cyclic Nucleotide Phosphodiesterase--Basic and Potential Clinical Applications, vol. 16, *Advances in Cyclic Nucleotide and Protein Phosphorylation Research*, pp. 245-259 (1984).
- Tulshian, D. et al., Synthesis and Phosphodiesterase Activity of Carboxylic Acid Mimetics of Cyclic Guanosine 3',5'-Monophosphate, *J. Med. Chem.* vol. 36, 1210-1220 (1993).
- Yasumoto, T. et al., Properties of Base-Substituted and Carboxyl-Esterified Analogues of Griseolic Acid, a Potent cAMP Phosphodiesterase Inhibitor, *Biochemical Pharmacology*, vol. 43, No. 10, pp. 2073,2081 (1992).
- Broughton, B.J. et al., Antiallergic Activity of 2-Phenyl-8-azaprin-6-ones, *Journal of Medicinal Chemistry*, vol. 18, No. 11, pp. 1117-1118 (1975).
- Kodama, K. et al., Effects of a novel, selective and potent phosphodiesterase type V inhibitor, E4021, on myocardial ischemia in guinea pigs, *Euro. J. of Pharma.* 263, pp. 93-99 (1994).
- Zacharski, L. R. et al., Effect of Mopidamol on Survival in Carcinoma of the Lung and Colon: Final Report of Veterans Administration Cooperative Study No. 188, *J. of the Nat'l. Cancer Inst.*, vol. 80, No. 2, pp. 90-96 (1988).
- Lichtner, R. B. et al., The Pyrimido-pyrimidine Derivatives RA 233 and RX-RA 85 affect Growth and Cytoskeletal Organization of Rat Mammary Adenocarcinoma Cells, *Eur. J. Cancer Clin. Oncol.*, vol. 23, No. 9, pp. 1269-1275 (1987).
- Janik, P. et al., Inhibition of Growth of Primary and Metastatic Lewis Lung carcinoma Cells by the Phosphodiesterase Inhibitor Isobutylmethylxanthine, *Cancer Res.* vol. 40, pp. 1950-1954, (Jun. 1980).
- Bergstrand, Hakan et al., Effects of Antiallergic Agents, Compound 48/80, and Some Reference Inhibitors on the Activity of Partially Purified Human Lung Tissue Adenosine Cyclic 3',5'-Monophosphate and Guanosine Cyclic 3',5'-Monophosphate Phosphodiesterases. *Molecular Pharmacology*, 13, pp. 38-43 (1976).
- Drees, Markus et al., 3',5'-Cyclic Nucleotide Phosphodiesterase in Tumor Cells as Potential Target for Tumor Growth Inhibition, *Cancer Research* 53, pp. 3058-3061 (1993).
- Semmler, J. et al., Xanthine derivatives: comparison between suppression of tumor necrosis factor- α production and inhibition of cAMP phosphodiesterase activity, *Immunology* 78, pp. 520-525 (1993).
- Mehta, Rajendra et al., Structure-Activity Relationships of Brassinin in Preventing the Development of Carcinogen-Induced Mammary Lesions in Organ Culture, *Anticancer Research* 14: 1209-1214 (1994).
- Makaryan, A.P. et al., Cyclic Nucleotides in Patients with Malignant Neoplasms of the Colon, *Laboratory of DeLo*, vol. 8, pp. 31-33 (1991).
- Carter et al., *Chemotherapy of Cancer*, 2nd ed., John Wiley & Sons, NY, NY, 1981, pp. 362-365.
- Biddle, William et al., Antineoplastic Effect of the Pyrimido-Pyrimidine Derivative: RA 233, *Pathologie Biologie*, Jan., 1984, pp. 9-13.
- Clarke, W.R. et al., The type III phosphodiesterase inhibitor milrinone and type V PDE inhibitor dipyridamole individually and synergistically reduce elevated pulmonary vascular resistance Abstract Only, *Pulm. Pharmacol.*, 7(2), pp. 81-89, (1984).
- Raeburn, David et al., Effects of isoenzyme-selective inhibitors of cyclic nucleotide phosphodiesterase on microvascular leak in guinea pig airways in vivo (Abstract Only), *J. Pharmacol. Exp. Ther.*, 267(3), pp. 1147-1151 (1993).
- Marcoz, P. et al., Modulation of rat thymocyte proliferative response through the inhibition of different cyclic nucleotide phosphodiesterase isoforms by means of selective inhibitors and CGMP-elevating agents (Abstract Only), *Mol. Pharmacol.* 44(5)

pp. 1027-1035 (1993).

Barnett, Mary S. et al., Initial biochemical and functional characterization of cyclic nucleotide phosphodiesterase isozymes in canine colonic smooth muscle (Abstract Only), J. Pharmacol. Exp. Ther., 264(2) pp. 801-812 (1993).

Molnar-Kimber, K. et al., Modulation of TNF α and IL-1 β from indotoxin-stimulated monocytes by selective PDE isozyme inhibitors (Abstract Only), Agents Actions 39(Spec. Conf. Issue), C77-C79 (1993).

Giorgi, Mauro et al., Characterization of 3':5' cyclic nucleotide phosphodiesterase activities of mouse neuroblastoma N18TG2 cells (Abstract Only), FEBS Lett. 324(1) pp. 76-80 (1993).

Porter, Roderick et al., Preparation of 6-phenyl-3-(5-tetrazolyl)pyridin-2(H)-one derivatives as cyclic AMP-dependent protein kinase agonists (Abstract Only), PCT Int. Appl. WO9206085 A1, (Sep. 26, 1991).

Molnar-Kimber, K.L. et al., Differential regulation of TNF- α and IL-1 β production from endotoxin stimulated human monocytes by phosphodiesterase inhibitors (Abstract Only), Mediators Inflammation 1(6) pp. 411-417 (1992).

Radomski, Marek W. et al., Human Colorectal adenocarcinoma cells; differential nitric oxide synthesis determines their ability of aggregate platelets (Abstract Only), Cancer Res. 51(22) pp. 6073-6078 (1991).

Anderson, Thomas L. G. et al., Interactions between isoprenaline, sodium nitroprusside, and isozyme-selective phosphodiesterase inhibitors on ADP-induced aggregation and cyclic Nucleotide levels in human platelets (Abstract Only), J. Cardiovasc. Pharmacol. 18(2) pp. 237-242 (1991).

Souness, John E. et al., Role of Selective cyclic GMP phosphodiesterase inhibition in the myorelaxant actions of M&B 22,943, MY-5445, vinpocetine and 1-methyl-3-isobutyl-8-(methylamino)xanthine (Abstract Only), Br. J. Pharmacol. 98(3) pp. 725-734 (1989).

Lichtner, Rosemarie B., The pyrimidopyrimidine derivatives RA233 and RX-RA85 affect cell cycle distribution of two murine tumor cell lines (Abstract Only), Eur. J. Cancer Clin. Oncol. 25(6), pp. 945-951 (1989).

Mamytbekova, A., et al., Antimetastatic effect of flurbiprofen and other platelet aggregation inhibitors (Abstract Only), Neoplasma 33(4), pp. 417-421 (1986).

Hagiwara, Masatoshi et al., Effect of 1-(3-chloroanilino)-4-phenylphthalazine (MY-5445), a specific inhibitor of cyclic CMP phosphodiesterase, on human platelet aggregation (Abstract Only), J. Pharmacol. Exp. Ther. 229(2) pp. 467-471 (1984).

ART-UNIT: 164

PRIMARY-EXAMINER: Goldberg; Jerome D.

ATTY-AGENT-FIRM: Stevenson; Robert W.

ABSTRACT:

A method for inhibiting neoplastic cells and related conditions by exposing them to substituted N-arylmethyl and heterocyclmethyl-1H-pyrazolo[3,4-B]quinolin-4-amines.

15 Claims, 0 Drawing figures